Pt. 40, App. B

d. Must be individually wrapped in a sealed plastic bag or shrink wrapping; or must have a peelable, sealed lid or other easily visible tamper-evident system.

e. May be made available separately at collection sites to address shy bladder situations when several voids may be required to complete the testing process.

2. Plastic Specimen Bottles

a. Each bottle must be large enough to hold at least 35 mL; or alternatively, they may be two distinct sizes of specimen bottles provided that the bottle designed to hold the primary specimen holds at least 35 mL of urine and the bottle designed to hold the split specimen holds at least 20 mL.

b. Must have screw-on or snap-on caps that prevent seepage of the urine from the bottles during shipment.

c. Must have markings clearly indicating the appropriate levels (30 mL for the primary specimen and 15 mL for the split) of urine that must be poured into the bottles.

d. Must be designed so that the required tamper-evident bottle seals made available on the CCF fit with no damage to the seal when the employee initials it nor with the chance that the seal overlap would conceal printed information.

e. Must be wrapped (with caps) together in a sealed plastic bag or shrink wrapping separate from the collection container; or must have peelable, sealed lid or other easily visible tamper-evident system.

3. Leak-Resistant Plastic Bag

a. Must have two sealable compartments or pouches which are leak-resistant; one large enough to hold two specimen bottles and the other large enough to hold the CCF paperwork.

b. The sealing methodology must be such that once the compartments are sealed, any tampering or attempts to open either compartment will be evident.

4. Absorbent material

Each kit must contain enough absorbent material to absorb the entire contents of both specimen bottles. Absorbent material must be designed to fit inside the leak-resistant plastic bag pouch into which the specimen bottles are placed.

5. Shipping Container

a. Must be designed to adequately protect the specimen bottles from shipment damage in the report of specimens from the collection site to the laboratory (e.g., standard courier box, small cardboard box, plastic container).

b. May be made available separately at collection sites rather than being part of an actual kit sent to collection sites.

c. A shipping container is not necessary if a laboratory courier hand-delivers the specimen bottles in the plastic leak-proof bags from the collection site to the laboratory.

APPENDIX B TO PART 40—DOT DRUG TESTING SEMI-ANNUAL LABORATORY REPORT TO EMPLOYERS

The following items are required on each laboratory report:

1. Specimen Results Reported (total number)
   By Test Reason
   (a) Pre-employment (number)
   (b) Post-Accident (number)
   (c) Random (number)
   (d) Reasonable Suspicion/Cause (number)
   (e) Return-to-Duty (number)
   (f) Follow-up (number)
   (g) Type of Test Not Noted on CCF (number)

2. Specimens Reported
   (a) Negative (number)
   (b) Negative and Dilute (number)

3. Specimens Reported as Rejected for Testing (total number)
   By Reason
   (a) Fatal flaw (number)
   (b) Uncorrected Flaw (number)

4. Specimens Reported as Positive (total number) By Drug
   (a) Marijuana Metabolite (number)
   (b) Cocaine Metabolite (number)
   (c) Opiates (number)
      (1) Codeine (number)
      (2) Morphine (number)
      (3) 6–AM (number)
   (d) Phencyclidine (number)
   (e) Amphetamines (number)
      (1) Amphetamine (number)
      (2) Methamphetamine (number)
      (3) MDMA (number)
      (4) MDA (number)
      (5) MDEA (number)

5. Adulterated (number)

6. Substituted (number)

7. Invalid Result (number)

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